## We claim:

10

15

20

25

35

1. A process for forming an extended polyalkylene-grafted interpolymer or gel, comprising:

a) mixing

ŧ.,

- 1) a polymer comprising mer units derived from maleic anhydride and mer units derived from at least one of
  - (A) a vinyl aromatic monomer, and
  - (B) an R¹(R²)ethylene monomer in which R¹ and R² independently are selected from H and substituted or unsubstituted C₁-C₂₀ alkyl or alkoxyl groups, and
- a maleated polyalkylene,
  so as to form a blend;
- b) to said blend, adding a dramine and allowing it to react with the mer units derived from maleric anhydride and with the maleated polyalkylene to form a polyalkylene grafted interpolymer; and
- allowing the polyalkylene-grafted interpolymer to cool in said mixer and adding an extender to provide said extended polyalkylene-grafted interpolymer; and
- d) optionally, extruding the extended polyalkylene-grafted interpolymer to form a get having a tan  $\delta$  of at least 0.3.
- 2. The process of claim 1 wherein step a) includes from about 50 to about 99 weight % of said polymer and from about 1 to about 50 weight % of said maleated polyalkylene and wherein step b) includes from about 0.1 to about 10 weight % of said diamine.
  - 3. The process of any of claims 1 to 2 wherein said polymer comprises from about 0.01 to about 5 weight percent mer units derived from maleic anhydride.
- 4. The process of any of claims 1 to 3 wherein steps b), c) and, optionally a) are carried out sequentially in a mixer, with no physical manipulation of said polyalkylene grafted interpolymer prior to step c) and without removing any of the materials from said mixer.



5

10

15

20

25

30

35

- 5. The process of any of claims 1 to 4 wherein step c) includes mixing about 10 weight percent to about 90 weight percent of said polyalkylene-grafted interpolymer with a total amount of about 90 weight percent to about 10 weight percent of said extender.
- 6. The process of any of claims 1 to 5 wherein at least one of the following is true:

said vinyl aromatic monomer is selected from any one or more of styrene, α-methylstyrene, ρ-methylstyrene, 4-phenylstyrene, m-methylstyrene, ο-methylstyrene, ρ-tert-butylstyrene, and dimethylstyrene.

R<sup>1</sup> and R<sup>2</sup> independently are selected from methyl, ethyl, propyl, isopropyl, butyl, isobutyl, pentyl, isopentyl, hexyl, heptyl, octyl, nofyl, decyl, undecyl, dodecyl, tridecyl, tetradecyl, pentadecyl, hexadecyl, heptadecyl, octadecyl, nonadecyl, eicosyl, cyclopropyl, 2,2-dimethylcyclopropyl, cyclopentyl, cyclohexyl, methoxymethyl, methoxyethyl, methoxypropyl, methoxybutyl, methoxypentyl, methoxyhexyl, methoxyoctyl, methoxynonyl, ethoxydecyl, ethoxymethyl, ethoxyethyl, ethoxypropyl, ethoxybutyl, ethoxypentyl, ethoxyhexyl, ethoxyheptyl, ethoxyoctyl, ethoxynonyl, ethoxydecyl, propoxymethyl, propoxyethyl, propoxypropyl, propoxybutyl, propoxypentyl, propoxyheptyl, propoxyoctyl, propoxynonyl, propoxydecyl, butoxymethyl, butoxyethyl, butoxypropyl, butoxybutyl, butoxypentyl, butoxyhexyl, butoxyhéptyl, butoxyoctyl, butoxynonyl, butoxydecyl, pentyloxymethyl, pentyloxyethyl, pentyloxypropyl, pentyloxybutyl, pentyloxypentyl, pentyoxyhexyl, pentyloxyoctyl, pentyloxynonyl, pentyloxydecyl, hexyloxymethyl, hexyloxyethyl, hexyloxyprop/l, hexyloxybutyl, hexyloxypentyl, hexyloxyhexyl, hexyloxyheptyl, hexyloxygctyl, hexyloxynonyl, hexyloxydecyl, heptyloxymethyl, heptyloxyethyl, heptyloxypropyl, heptyloxybutyl, hexyloxypentyl, heptyloxyhexyl, heptyloxyheptyl, heptyloxyoctyl, heptyloxynonyl, heptyloxydecyl, octyloxymethyl, oxtyloxyethyl, octyloxypropyl, octyloxybutyl, octyloxypentyl, octyloxyhexyl, octyloxyheptyl, ogtyloxyoctyl, oxtyloxynonyl, decyloxymethyl, decyloxyethyl, decyloxypropyl/decyloxybutyl, decyloxypentyl, decyloxyhexyl, decyloxyheptyl, 1methylethyl, 1/methylpropyl, 1-methylbutyl, 1-methylpentyl, 1methylhepty/I, 1-methyloctyl, 1-methylnonyl, 1-methyldecyl, 2-methylpropyl, 2methylbyfyl, 2-methylpentyl, 2-methylhexyl, 2-methylheptyl, 2-methyloctyl, 2,3,3trimethy/butyl, 3-methylpentyl, 2,3-dimethylpentyl, 2,4-dimethylpentyl, 2,3,3,4tetramethylpentyl, 3-methylhexyl, and 2,5-dimethylhexyl.

5

10

15

25

- 7. The process of any of claims 1 to 6 wherein said maleated polyalkylene includes mer units derived from at least one of ethylene and propylene.
- 8. The process of any of claims 1 to 7 wherein, prior to step a), an amine is reacted with a portion of the mer units derived from the maleic anhydride to form maleimide mer units, wherein at least one of the following optionally is true of said imidization:

it is carried out immediately preceding step a) with no physical manipulation of the polyn.er prior to step a), and

it and step a) are carried out in a mixer without removing the imidization product from the mixer.

- 9. The process of claim 8, wherein said amine comprises from 1 to 50 carbon atoms and is selected from at least one of alkyl amines, alkyl benzyl amines, alkyl phenyl amines, alkoxybenzyl amines, alkyl aminobenzoates, and alkoxy aniline.
  - 10. The process of claim 8, wherein said amine is a primary amine.

11. The process of any of claims 1 to 8 wherein said diamine is selected from aliphatic or cycloaliphatic diamines corresponding to the general formula R³(NH₂)₂ wherein R³ represents

1) a C<sub>2</sub> - C<sub>20</sub> aliphatic hydrocarbon group,

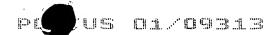
2) a C<sub>4</sub> - C<sub>20</sub> cycloalighatic hydrocarbon group,

3) a C<sub>6</sub> - C<sub>20</sub> aromátic hydrocarbon group, or

4) a C<sub>4</sub> - C<sub>20</sub> N/heterocyclic ring.

12. The process of claim 11, wherein said diamine is selected from at least one of ethylene diamine; 1,2-propylene diamine; 1,3-propylene diamine; 1,4-diaminobutane; 2,2-dimethyl-1,3-diaminopropane; 1,6-diaminohexane; 2,5-dimethyl-2,5-diaminohexane; 1,6-diamino-2,2,4-trimethyldiaminohexane; 1,8-diaminooctane; 1,10-diaminodecane; 1,11-diaminoundecane; 1,12-diaminododecane; 1-methyl-4-(aminoisopropyl)-cyclohexylamine; 3-aminomethyl-3,5,5-trimethyl-cyclohexylamine; 1,2-bis-(aminomethyl)-cyclobutane; 1,2-diamino-3,6-dimethylbenzene; 1,2-diaminocyclohexane; 1,4-diaminodecalin; 1,5-diaminodecalin; 1,8-diaminodecalin; 1-





methyl-4-aminoisopropyl-cyclohexylamine; 4,4'-diamino-dicyclohexyl; 4,4'-diamino-dicyclohexyl methane; 2,2'-(bis-4-amino-cyclohexyl)-propane; 3,3'-dimethyl-4,4'-diamino-dicyclohexyl methane; 1,2-bis-(4-aminocyclohexyl)-ethane; 3,3',5,5'-tetramethyl-bis-(4-aminocyclohexyl)-methane; 3,3',5,5'-tetramethyl-bis-(4-aminocyclohexyl)-propane; 1,4-bis-(2-aminoethyl)-benzene; benzidine; 4,4'-thiodianiline, 3,3'-dimethoxybenzidine; 2,4-diaminotoluene, diaminoditolylsulfone; 2,6-diaminopyridine; 4-methoxy-6-methyl-*m*-phenylenediamine; diaminodiphenyl ether; 4,4'-bis(o-toluidine); o-phenylenediamine; methylene-bis(o-chloroaniline); bis(3,4-diaminophenyl)sulfone; diaminodiphenylsulfone; 4-chloro-o-phenylenediamine; *m*-aminobenzylamine; *m*-phenylenediamine; 4,4'-C<sub>1</sub>-C<sub>6</sub>-dianiline; aniline-formaldehyde resin; trimethylene glycol di-p-aminobenzoate; bis-(2-aminoethyl)-amine; bis-(3-aminopropyl)-amine; bis-(4-aminobutyl)amine; bis-(6-aminohexyl)-amine; and isomeric mixtures of dipropylene triamine and dibutylene triamine.

y includ

5

10

15

20

13. A single batch process for preparing a polymer composition which includes a polyalkylene-grafted interpolymer, said process comprising:

- a) forming a maleimide interpolymer in a mixing vessel by reacting an amine with a portion of maleic anhydride-derived mer units of an interpolymer comprising maleic anhydride-derived mer units and at least one of
  - 1) vinyl aromatic-derived mer units, and
  - 2) R<sup>1</sup>(R<sup>2</sup>)ethylene-derived mer units in which R<sup>1</sup> and R<sup>2</sup> independently are H or substituted or unsubstituted C<sub>1</sub> to C<sub>20</sub> alkyl groups or alkoxyl groups;

 adding sufficient maleaded polyalkylene such that the mixing vessel contains from about 1 to about 50 weight percent maleated polyalkylene and from about 50 to about 99 weight percent maleimide interpolymer;

c) mixing from about 0.1 to about 10 weight % of a diamine with the maleimide interpolymer and maleated polyalkylene in the mixing vessel to form/said polyalkylene-grafted interpolymer; and

d) optionally, without removing the product of step c) from the vessel, cooling the polyalkylene-grafted interpolymer in the mixer to a temperature at which an extender is stable in the polyalkylene-grafted interpolymer, and adding an extender to the mixer.

25

30

35

28